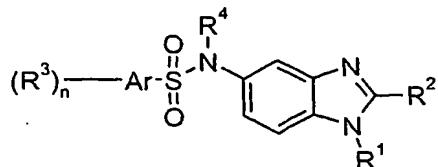


What is claimed is:

1. A compound of Formula I or a pharmaceutically acceptable salt thereof:



5

I

wherein

R¹ is selected from C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, R⁵-C(=O)-O-C₁₋₆alkyl, R⁵R⁶N-C₁₋₆alkyl, R⁵O-C₁₋₆alkyl, R⁵C(=O)N(-R⁶)-C₁₋₆alkyl, R⁵R⁶NS(=O)₂-C₁₋₆alkyl, R⁵CS(=O)₂N(-R⁶)-C₁₋₆alkyl, R⁵R⁶NC(=O)N(-R⁷)-C₁₋₆alkyl,

10 R⁵R⁶NS(=O)₂N(R⁷)-C₁₋₆alkyl, C₆₋₁₀aryl-C₁₋₆alkyl, C₆₋₁₀aryl-C(=O)-C₁₋₆alkyl, C₃₋₁₀cycloalkyl-C₁₋₆alkyl, C₄₋₈cycloalkenyl-C₁₋₆alkyl, C₃₋₆heterocyclyl-C₁₋₆alkyl, C₃₋₆heterocyclyl-C(=O)-C₁₋₆alkyl, C₁₋₁₀hydrocarbyl amino, R⁵R⁶N-, R⁵O-, R⁵C(=O)N(-R⁶)-, R⁵R⁶NS(=O)₂-, R⁵CS(=O)₂N(-R⁶)-, R⁵R⁶NC(=O)N(-R⁷)-, R⁵R⁶NS(=O)₂N(R⁷)-, C₆₋₁₀aryl, C₆₋₁₀aryl-C(=O)-, C₃₋₁₀cycloalkyl, C₄₋₈cycloalkenyl, C₃₋₆heterocyclyl and C₃₋₆heterocyclyl-C(=O)-; wherein said C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₆₋₁₀aryl-C₁₋₆alkyl, C₆₋₁₀aryl-C(=O)-C₁₋₆alkyl, C₃₋₁₀cycloalkyl-C₁₋₆alkyl, C₄₋₈cycloalkenyl-C₁₋₆alkyl, C₃₋₆heterocyclyl-C(=O)-C₁₋₆alkyl, C₁₋₁₀hydrocarbyl amino, C₆₋₁₀aryl, C₆₋₁₀aryl-C(=O)-, C₃₋₁₀cycloalkyl, C₄₋₈cycloalkenyl, C₃₋₆heterocyclyl or C₃₋₆heterocyclyl-C(=O)- used in defining R¹ is optionally substituted by one or more groups selected from halogen, cyano, nitro, methoxy, ethoxy, methyl, ethyl, hydroxy, benzyl, and -NR⁵R⁶;

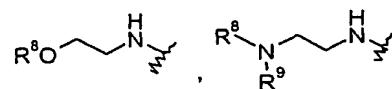
R² is selected from C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₆alkyl, C₄₋₈cycloalkenyl-C₁₋₆alkyl, C₃₋₆heterocycloalkyl-C₁₋₆alkyl, C₄₋₈cycloalkenyl, R⁵R⁶N-, C₃₋₅heteroaryl, C₆₋₁₀aryl and C₃₋₆heterocycloalkyl, wherein said C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₃₋₈cycloalkyl, C₃₋₈cycloalkyl-C₁₋₆alkyl, C₄₋₈cycloalkenyl-C₁₋₆alkyl, C₃₋₆heterocycloalkyl-C₁₋₆alkyl, C₄₋₈cycloalkenyl, C₃₋₆heteroaryl, C₆₋₁₀aryl or C₃₋₆heterocycloalkyl used in defining R² is optionally substituted by one or more groups selected from halogen, cyano, nitro, methoxy, ethoxy, methyl, ethyl, hydroxy, and -NR⁵R⁶;

wherein R⁵, R⁶ and R⁷ are independently selected from -H, C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, and a divalent C₁₋₆group that together with another divalent R⁵, R⁶ or R⁷ forms a portion of a ring;

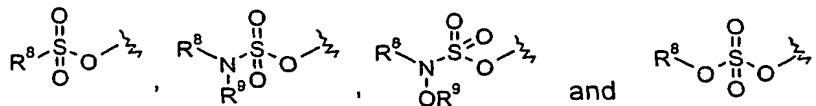
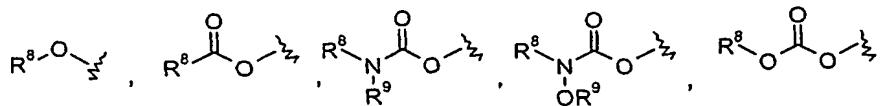
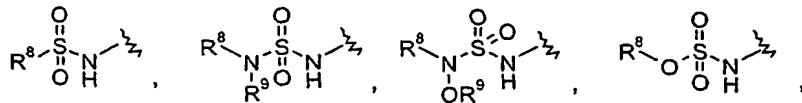
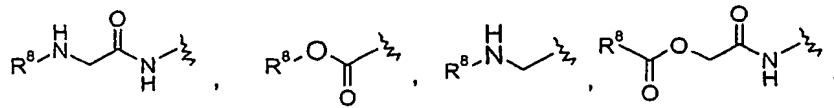
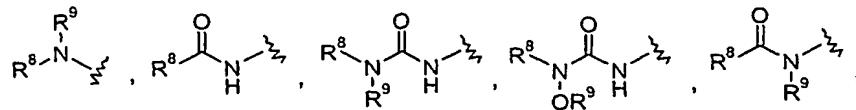
Ar is selected from C₆₋₁₀aryl and C₃₋₈heteroaryl;

5 n is selected from 0, 1, 2 and 3;

each of R³ is independently selected from -H, nitro, halogen, C₁₋₁₀alkyl C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₆alkyl, C₄₋₈cycloalkenyl-C₁₋₆alkyl, C₃₋₆heterocycloalkyl-C₁₋₆alkyl, C₃₋₆heterocycloalkyl and

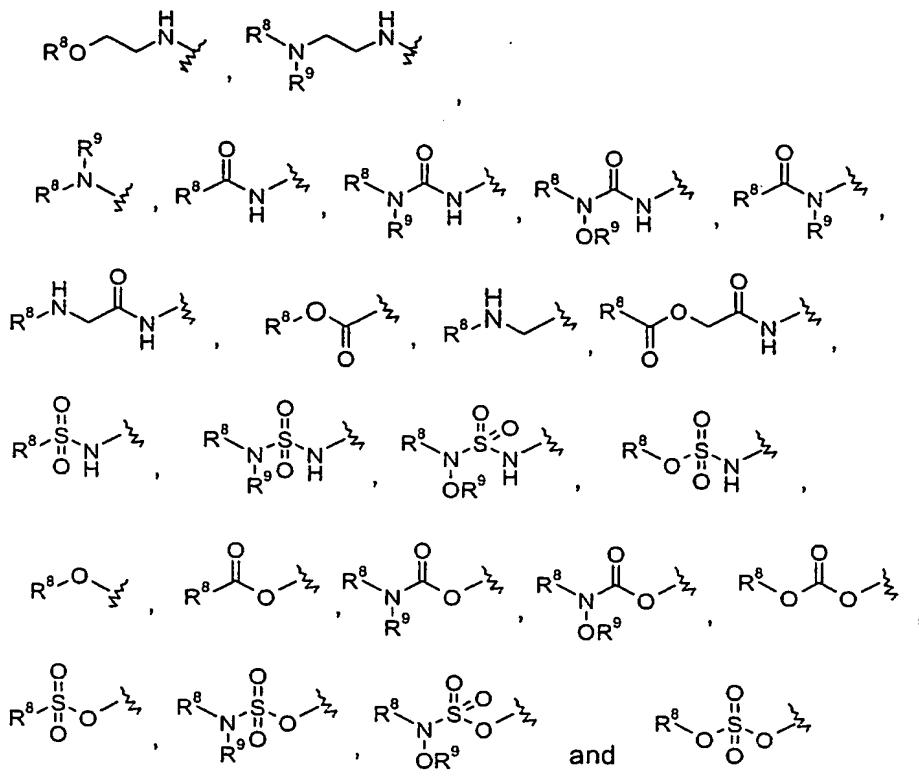


,



and

10 optionally substituted with one or more groups selected from C₁₋₆alkyl, hydroxy, halogen, amino and C₁₋₆alkoxy,



each of R⁸ and R⁹ is independently selected from -H, C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₆alkyl, C₃₋₆heterocyclyl, C₆₋₁₀aryl, C₃₋₆heterocyclyl-C₁₋₆alkyl, C₆₋₁₀aryl-C₁₋₆alkyl, and a divalent C₁₋₆group that together with another divalent group selected from R⁸ and R⁹ forms a portion of a ring,
 wherein said C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₆alkyl, C₃₋₆heterocyclyl-C₁₋₆alkyl, C₆₋₁₀aryl, C₃₋₆heterocyclyl-C₁₋₆alkyl, C₆₋₁₀aryl-C₁₋₆alkyl, or divalent C₁₋₆group is optionally substituted by one or more groups selected from halogen, cyano, nitro, methoxy, ethoxy, methyl, ethyl, hydroxy, and -NR⁵R⁶; and
 10 R⁴ is selected from -H, C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₆alkyl, and C₄₋₈cycloalkenyl-C₁₋₆alkyl.
 2. A compound as claimed in claim 1, wherein
 R¹ is selected from C₁₋₆alkyl, C₁₋₆alkyl-C(=O)-O-C₁₋₄alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, phenyl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl-C₁₋₄alkyl, C₄₋₆cycloalkenyl-C₁₋₄alkyl, C₃₋₆heterocyclyl-C₁₋₄alkyl, C₆₋₁₀aryl, C₃₋₆heterocyclyl, C₃₋₁₀cycloalkyl, and C₄₋₆cycloalkenyl, wherein said C₁₋₆alkyl, C₁₋₆alkyl-C(=O)-O-C₁₋₄alkyl, C₂₋₆alkenyl, C₂₋₆

6alkynyl, phenyl-C₁-alkyl, C₃-₁₀cycloalkyl-C₁-alkyl, C₄-₆cycloalkenyl-C₁-alkyl, C₆-₁₀aryl, C₃-₆heterocyclyl-C₁-alkyl, C₃-₆heterocyclyl, C₃-₁₀cycloalkyl, and C₄.

6cycloalkenyl used in defining R¹ is optionally substituted by one or more groups selected from halogen, cyano, nitro, methoxy, ethoxy, methyl, ethyl, hydroxy, benzyl, 5 and -NR⁵R⁶;

R² is selected from C₁-alkyl, C₂-alkenyl, C₃-₆cycloalkyl, C₃-₆cycloalkyl-C₁-alkyl, C₄-₆cycloalkenyl-C₁-alkyl, C₃-₆heterocycloalkyl-C₁-alkyl, C₄-₆cycloalkenyl, C₃-₅heteroaryl, R⁵R⁶N-, and phenyl, wherein said C₁-alkyl, C₂-alkenyl, C₃-₆cycloalkyl, C₃-₆cycloalkyl-C₁-alkyl, C₄-₆cycloalkenyl-C₁-alkyl,

10 C₃-₆heterocycloalkyl-C₁-alkyl, C₄-₆cycloalkenyl, C₃-₅heteroaryl, R⁵R⁶N-, and phenyl used in defining R² is optionally substituted by one or more groups selected from halogen, cyano, nitro, methoxy, ethoxy, methyl, ethyl, hydroxy and amino;

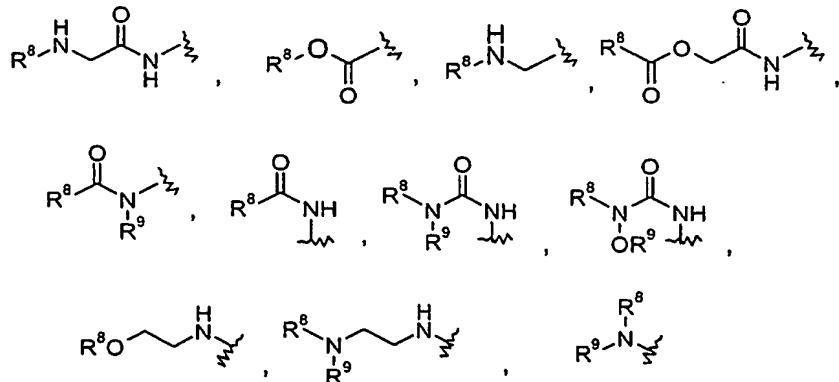
wherein R⁵ and R⁶ are independently selected from -H, C₁-alkyl, C₂-alkenyl, and a divalent C₁-alkylene that together with another divalent R⁵ or R⁶ and optionally 15 a heteroatom forms a portion of a ring;

Ar is selected from phenyl and C₃-₅heteroaryl;

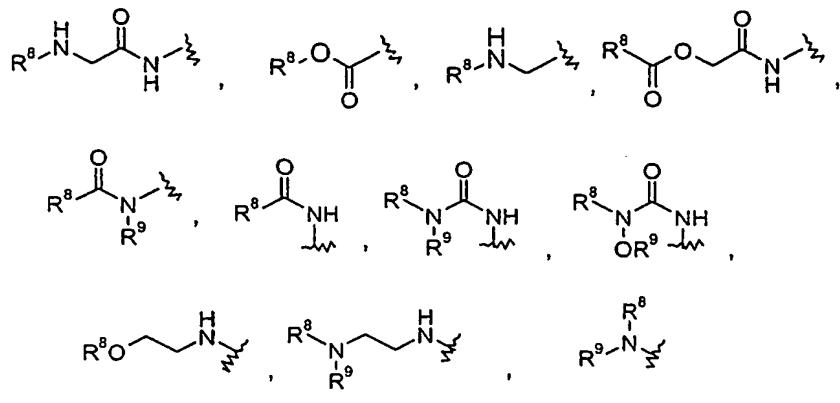
n is selected from 0, 1 and 2;

each of R³ is independently selected from -H, nitro, halogen, C₁-alkyl C₂-alkenyl, C₃-₆cycloalkyl, C₃-₆heterocycloalkyl-C₁-alkyl,

20



and, C₃-₆heterocycloalkyl optionally substituted with one or more groups selected from C₁-alkyl, hydroxy, halogen and



each of R⁸ and R⁹ is independently selected from -H, C₁₋₆alkyl, C₂₋₆alkenyl,

C₃₋₆cycloalkyl, C₃₋₆cycloalkyl-C₁₋₆alkyl, C₃₋₆heterocyclyl and C₃₋₆heterocyclyl-C₁₋₆alkyl, wherein said C₁₋₆alkyl, C₂₋₆alkenyl, C₃₋₆cycloalkyl, C₃₋₆cycloalkyl-C₁₋₆alkyl,

5 C₃₋₆heterocyclyl and C₃₋₆heterocyclyl-C₁₋₆alkyl are optionally substituted by one or more groups selected from halogen, cyano, nitro, methoxy, ethoxy, methyl, ethyl, hydroxy and -NR¹⁰R¹¹; and

R⁴, R¹⁰ and R¹¹ are independently selected from -H and C₁₋₃alkyl.

10 3. A compound as claimed claim 1,

wherein R¹ is selected from C₁₋₆alkyl, C₁₋₃alkyl-C(=O)-O-C₁₋₃alkyl,

C₂₋₆alkenyl, phenyl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl-C₁₋₄alkyl, C₄₋₆cycloalkenyl-C₁₋₄alkyl, C₃₋₆heterocylcoalkyl-C₁₋₄alkyl, C₆₋₁₀aryl, C₃₋₁₀cycloalkyl, and C₄₋₆cycloalkenyl, wherein said C₁₋₆alkyl, C₂₋₆alkenyl, phenyl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl-C₁₋₄alkyl,

15 C₄₋₆cycloalkenyl-C₁₋₄alkyl, C₃₋₆heterocylcoalkyl-C₁₋₄alkyl, C₆₋₁₀aryl, C₃₋₁₀cycloalkyl, and C₄₋₆cycloalkenyl used in defining R¹ is optionally substituted by one or more groups selected from halogen, methoxy, ethoxy, methyl, ethyl, hydroxy, benzyl, and amino;

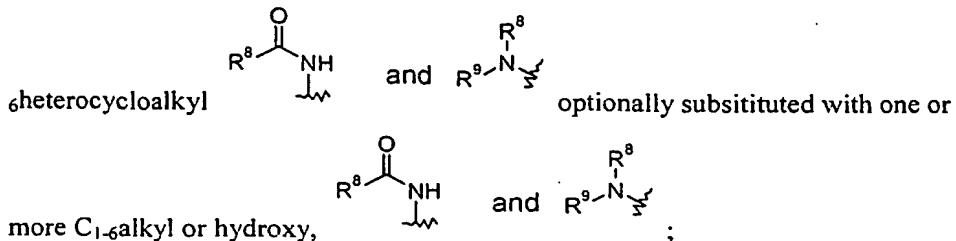
R² is selected from C₁₋₆alkyl, C₂₋₆alkenyl, C₃₋₆cycloalkyl and C₃₋₆cycloalkyl-

20 C₁₋₄alkyl, wherein said C₁₋₆alkyl, C₂₋₆alkenyl, C₃₋₆cycloalkyl and C₃₋₆cycloalkyl-C₁₋₄alkyl used in defining R² is optionally substituted by one or more groups selected from halogen, methoxy, ethoxy, methyl, ethyl, hydroxy and amino;

Ar is selected from phenyl and C₃₋₅heteroaryl and

n is selected from 0, 1 and 2;

each of R³ is independently selected from -H, halogen, nitro, C₁₋₃alkyl, C₃-



wherein said C₃₋₆heterocycloalkyl contain at least one nitrogen ring atom and

- 5 the radical of C₃₋₆heterocycloalkyl is located on the at least one nitrogen ring atom,
and wherein each of R⁸ and R⁹ is independently selected from -H, C₁₋₆alkyl,
morpholinyl- C₁₋₃alkyl, pyrrolidinyl-C₁₋₃alkyl, and piperidinyl-C₁₋₃alkyl, wherein said
C₁₋₆alkyl, morpholinyl- C₁₋₃alkyl, pyrrolidinyl-C₁₋₃alkyl, and piperidinyl-C₁₋₃alkyl are
optionally substituted by one or more groups selected from halogen, methoxy, ethoxy,
10 methyl, ethyl, hydroxy and -NR⁵R⁶; and

R⁴, R⁵ and R⁶ are independently selected from -H and C₁₋₃alkyl.

4. A compound as claimed in claim 1, wherein

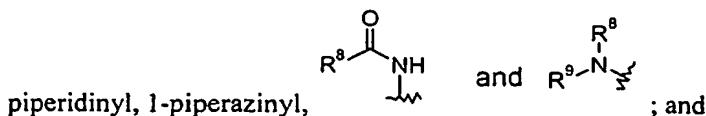
- R¹ is selected from cyclohexylmethyl, cyclopentylmethyl, cyclobutylmethyl,
15 cyclopropylmethyl, cyclohexylethyl, cyclopentylethyl, bicyclo[2.2.1]hept-5-en-2-
ylmethyl, 4,4-difluorocyclohexylmethyl, tetrahydropyranylmethyl,
tetrahydropyranylethyl, tetrahydrofuranylmethyl, 1-piperidinylethyl, and N-methyl-2-
piperidinylmethyl;

- 20 R² is selected from t-butyl, n-butyl, 2-methyl-2-butyl, isopentyl, 2-methoxy-2-
propyl, 2-hydroxyl-propyl, trifluoromethyl, 1,1-difluoroethyl, 2,2,2-trifluoroethyl, 1-
methyl-propyl, 1,1-dimethyl-propyl, 1,1-dimethyl-3-buten-1-yl, ethyl, and 2-propyl;

Ar is selected from phenyl, pyridyl, pyrimidyl, thiazolyl, thienyl, isoxazolyl,
imidazolyl, and pyrazolyl;

n is selected from 0, 1 and 2;

- 25 each of R³ is independently selected from -H, C₁₋₃alkyl, 4-morpholinyl, 1-



wherein 4-morpholinyl, 1-piperidinyl, and 1-piperazinyl are optionally substituted with one or more methyl; and wherein

each of R⁸ and R⁹ is independently selected from –H, C₁₋₃alkyl, morpholinylmethyl, pyrrolidinyl-methyl, and piperidinyl-methyl, wherein said

5 C₁₋₃alkyl, morpholinylmethyl, pyrrolidinyl-methyl, and piperidinyl-methyl are optionally substituted by one or more groups selected from hydroxy, amino and dimethylamino.

5. A compound selected from:

10 N-[2-*tert*-Butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl]thiophene-2-sulfonamide;
N-[2-*tert*-Butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl]-N-methylthiophene-2-sulfonamide;
N-(1-Benzyl-2-*tert*-butyl-1*H*-benzimidazol-5-yl)-N-methylbenzenesulfonamide;

15 N-[2-*tert*-Butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl]-N,3,5-trimethylisoxazole-4-sulfonamide;
N-[2-*tert*-Butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl]-N,1,2-trimethyl-1*H*-imidazole-4-sulfonamide;
N-[2-*tert*-Butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl]-N,1,3,5-tetramethyl-1*H*-pyrazole-4-sulfonamide;
20 N-[2-*tert*-butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl]benzene sulphonamide;
N-[1-(cyclohexylmethyl)-2-ethyl-1*H*-benzimidazol-5-yl]benzenesulfonamide;
N-[1-(cyclohexylmethyl)-2-isopropyl-1*H*-benzimidazol-5-yl]benzene
25 sulphonamide;
N-[1-(cyclohexylmethyl)-2-(1-methylcyclopropyl)-1*H*-benzimidazol-5-yl]benzenesulfonamide;
N-[1-(cyclohexylmethyl)-2-(1,1-dimethylpropyl)-1*H*-benzimidazol-5-yl]-benzenesulfonamide;

30 N-[1-(cyclohexylmethyl)-2-(1,1-dimethyl-3-butenyl)-1*H*-benzimidazol-5-yl]-benzenesulfonamide;
N-[1-(cyclohexylmethyl)-2-(1-methyl-4-piperidinyl)-1*H*-benzimidazol-5-yl]-benzenesulfonamide;

N-[1-(cyclohexylmethyl)-2-(1,1-dimethylethyl)-1*H*-benzimidazol-5-yl]-*N*-methylbenzenesulfonamide;

N-[1-(cyclohexylmethyl)-2-ethyl-1*H*-benzimidazol-5-yl]-*N*-methyl-benzene sulphonamide;

5 *N*-[1-(cyclohexylmethyl)-2-isopropyl-1*H*-benzimidazol-5-yl]-*N*-methyl-benzene sulphonamide;

N-[1-(cyclohexylmethyl)-2-(1-methylcyclopropyl)-1*H*-benzimidazol-5-yl]-*N*-methyl-benzenesulfonamide;

10 *N*-[1-(cyclohexylmethyl)-2-(1-methyl-4-piperidinyl)-1*H*-benzimidazol-5-yl]-*N*-methyl- benzenesulfonamide;

 4-[1-(cyclohexylmethyl)-5-[methyl(phenylsulfonyl)amino]-1*H*-benzimidazol-2-yl]-1,1-dimethyl- piperidinium;

N-[2-(1,1-dimethylethyl)-1-[(tetrahydro-2*H*-pyran-4-yl)methyl]-1*H*-benzimidazol-5-yl]-benzenesulfonamide;

15 *N*-[2-(1,1-dimethylethyl)-1-[(tetrahydro-2-furanyl)methyl]-1*H*-benzimidazol-5-yl]-benzenesulfonamide;

N-[1-(cyclobutylmethyl)-2-(1,1-dimethylethyl)-1*H*-benzimidazol-5-yl]-benzenesulfonamide;

20 *N*-[1-(cyclopropylmethyl)-2-(1,1-dimethylethyl)-1*H*-benzimidazol-5-yl]-benzenesulfonamide;

N-(4-{[[2-*tert*-butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl) acetamide;

N-[2-*tert*-Butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl]-*N*-methyl-6-morpholin-4-ylpyridine-3-sulfonamide;

25 *N*-[2-*tert*-Butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl]-*N*-methyl-4-nitrobenzenesulfonamide;

 4-Amino-*N*-[2-*tert*-butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl]-*N*-methylbenzenesulfonamide;

30 *N*-(4-{[[2-*tert*-Butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)propanamide;

N-(4-{[[2-*tert*-Butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)-2-methylpropanamide;

N-(4-{{[2-*tert*-Butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)-2,2-dimethylpropanamide;

N-[2-*tert*-Butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl]-4-(ethylamino)-*N*-methylbenzenesulfonamide;

5 *N*-[2-*tert*-Butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl]-4-(formylamino)-*N*-methylbenzenesulfonamide;

2-Bromo-*N*-(4-{{[2-*tert*-butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)acetamide;

10 *N*-(4-{{[2-*tert*-Butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)-2-pyrrolidin-1-ylacetamide;

*N*¹-(4-{{[2-*tert*-Butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)-*N*²,*N*²-dimethylglycinamide;

N-(4-{{[2-*tert*-Butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)-2-morpholin-4-ylacetamide;

15 *N*¹-(4-{{[2-*tert*-Butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)glycinamide;

 2-[{(4-{{[2-*tert*-Butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)amino]-2-oxoethyl acetate;

N-(4-{{[2-*tert*-Butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)-2-hydroxyacetamide;

20 *N*-[1-(cyclohexylmethyl)-2-(1,1-dimethylethyl)-1*H*-benzimidazol-5-yl]-*N*-methyl-4-(4-morpholinyl)-benzenesulfonamide;

N-[1-(cyclohexylmethyl)-2-(1,1-dimethylethyl)-1*H*-benzimidazol-5-yl]-*N*-methyl-4-(4-methyl-1-piperazinyl)-benzenesulfonamide;

25 *N*-[2-*tert*-Butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]-*N*-methylbenzenesulfonamide;

N-[2-*tert*-Butyl-1-(tetrahydro-2*H*-pyran-2-ylmethyl)-1*H*-benzimidazol-5-yl]-*N*-methylbenzenesulfonamide;

N-[1-(cyclohexylmethyl)-2-(1-hydroxy-1-methylethyl)-1*H*-benzimidazol-5-yl]-benzenesulfonamide;

30 *N*-[1-(cyclohexylmethyl)-2-(1-methoxy-1-methylethyl)-1*H*-benzimidazol-5-yl]-*N*-methyl-benzenesulfonamide;

N-[1-(cyclohexylmethyl)-2-(1-methoxy-1-methylethyl)-1*H*-benzimidazol-5-yl]—
benzenesulfonamide;

N-[2-*tert*-Butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]-
N,1,2-trimethyl-1*H*-imidazole-5-sulfonamide;

5 Ethyl 4-{{[2-*tert*-butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-
yl](methyl)amino}sulfonyl}-3,5-dimethyl-1*H*-pyrrole-2-carboxylate;

N-[2-*tert*-Butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]-4-
(hydroxymethyl)-*N*-methylbenzenesulfonamide;

10 *N*-[2-*tert*-Butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]-*N*-
methyl-4-(1*H*-1,2,3-triazol-1-ylmethyl)benzenesulfonamide;

N-[2-*tert*-Butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]-4-
{[(2-hydroxyethyl)amino]methyl}-*N*-methylbenzenesulfonamide;

15 *N*-[2-*tert*-Butyl-1-(cyclopentylmethyl)-1*H*-benzimidazol-5-yl]-*N*-
methylbenzenesulfonamide;

N-[2-*tert*-Butyl-1-(2-cyclohexylethyl)-1*H*-benzimidazol-5-yl]-*N*-
methylbenzenesulfonamide;

N-[1-(1-Benzylpyrrolidin-3-yl)-2-*tert*-butyl-1*H*-benzimidazol-5-yl]-*N*-
methylbenzenesulfonamide;

20 *N*-{2-*tert*-Butyl-1-[(4,4-difluorocyclohexyl)methyl]-1*H*-benzimidazol-5-yl}-*N*-
methylbenzenesulfonamide;

N-[2-*tert*-Butyl-1-(pyridin-4-ylmethyl)-1*H*-benzimidazol-5-yl]-*N*-
methylbenzenesulfonamide;

25 *N*-methyl-*N*-[1-(tetrahydro-2*H*-pyran-4-ylmethyl)-2-(trifluoromethyl)-1*H*-
benzimidazol-5-yl]benzenesulfonamide;

N-[2-(1,1-difluoroethyl)-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-
yl]-*N*-methylbenzenesulfonamide;

N-methyl-*N*-[1-(tetrahydro-2*H*-pyran-4-ylmethyl)-2-(2,2,2-trifluoroethyl)-1*H*-
benzimidazol-5-yl]benzenesulfonamide;

30 *N*-[1-(cyclohexylmethyl)-2-(1-ethylpropyl)-1*H*-benzimidazol-5-
yl]benzenesulfonamide;

N-[1-(cyclohexylmethyl)-2-(1-ethylpropyl)-1*H*-benzimidazol-5-yl]-*N*-
methylbenzenesulfonamide; *N*-[2-*tert*-butyl-1-(cyclohexylmethyl)-1*H*-
benzimidazol-5-yl]-*N*-ethylbenzenesulfonamide;

N-methyl-*N*-[2-(1-methyl-1-pyridin-2-ylethyl)-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]benzenesulfonamide;
N-[2-(1-cyano-1-methylethyl)-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]-*N*-methylbenzenesulfonamide;

5 *N*-methyl-*N*-[2-propyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]benzenesulfonamide;
 5-Bromo-*N*-[2-*tert*-butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl]-6-chloro-*N*-methylpyridine-3-sulfonamide;
 5-Bromo-*N*-[2-*tert*-butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl]-6-[(2-hydroxyethyl)amino]-*N*-methylpyridine-3-sulfonamide;

10 *N*-[2-*tert*-Butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl]-6-[(2-hydroxyethyl)amino]-*N*-methylpyridine-3-sulfonamide;
 N-(5-{{[2-*tert*-Butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}pyridin-2-yl)acetamide;

15 *N*-(3-{{[2-*tert*-Butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)acetamide;
 *N*¹-(4-{{[2-*tert*-Butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)-*N*²-(2-hydroxyethyl)glycinamide;
 4-[(Aminocarbonyl)amino]-*N*-[2-*tert*-butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]-*N*-methylbenzenesulfonamide;

20 *N*-(4-{{[2-*tert*-Butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)acetamide;
 N-(4-{{[2-*tert*-Butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)-*N*-methylacetamide;

25 *N*-(4-{{[2-*tert*-Butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)-2,2-dimethylpropanamide;
 N-(4-{{[2-*tert*-Butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)-2-hydroxyacetamide;
 *N*¹-(4-{{[2-*tert*-Butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)-*N*²,*N*²-dimethylglycinamide;

30 *N*¹-(4-{{[2-*tert*-Butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)glycinamide;

*N*¹-(4-{{[2-*tert*-Butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)-*N*²-methylglycinamide;
5 *N*-[2-*tert*-Butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]-6-[(2-hydroxyethyl)amino]-*N*-methylpyridine-3-sulfonamide;
 N-[2-*tert*-Butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]-6-[(2-methoxyethyl)amino]-*N*-methylpyridine-3-sulfonamide;
 N-[2-*tert*-Butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]-6-(formylamino)-*N*-methylpyridine-3-sulfonamide;
 N-(5-{{[2-*tert*-Butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}pyridin-2-yl)acetamide;
 N-[4-({{[2-*tert*-Butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]amino}sulfonyl}phenyl]acetamide;
 N-[4-({{[2-*tert*-Butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl]amino}sulfonyl}phenyl]acetamide;
15 *N*-(4-{{[2-*tert*-Butyl-1-(2-piperidin-1-ylethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)acetamide;
 N-(4-{{[2-*tert*-Butyl-1-(1,4-dioxan-2-ylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)acetamide;
 N-(4-{{[2-*tert*-Butyl-1-[(1-methylpiperidin-2-yl)methyl]-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)acetamide;
20 *N*-(4-{{[(2-*tert*-Butyl-1-[(2*R*)-1-methylpiperidin-2-yl]methyl}-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)acetamide;
 N-[4-({methyl}[1-(tetrahydro-2*H*-pyran-4-ylmethyl)-2-(trifluoromethyl)-1*H*-benzimidazol-5-yl]amino]sulfonyl)phenyl]acetamide;
25 4-Bromo-*N*-[1-(cyclohexylmethyl)-2-(1,1-dimethylethyl)-1*H*-benzimidazol-5-yl]-*N*-methyl-benzenesulfonamide;
 N-[2-*tert*-butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl]-4-[(2-hydroxyethyl)amino]-*N*-methylbenzenesulfonamide;
 N-[2-*tert*-butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl]-4-(dimethylamino)-
30 *N*-methylbenzenesulfonamide;
 4-[bis(2-hydroxyethyl)amino]-*N*-[2-*tert*-butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl]-*N*-methylbenzenesulfonamide;

*N-[2-*tert*-butyl-1-(cyclohexylmethyl)-1*H*-benzimidazol-5-yl]-N,4-dimethyl-3,4-dihydro-2*H*-1,4-benzoxazine-7-sulfonamide;*

*N-[4-{(methyl[2-(1-methyl-1-pyridin-2-ylethyl)-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]amino}sulfonyl]phenyl]acetamide;*

5 *N-(4-{[[2-*tert*-butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl](ethyl)amino}sulfonyl)phenyl]acetamide;*

*4-[(aminocarbonyl)amino]-N-[2-*tert*-butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]-N-ethylbenzenesulfonamide;*

10 *N-[2-*tert*-butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]-N-ethyl-4-{{(methylamino)carbonyl}amino}benzenesulfonamide;*

*4-amino-N-[2-*tert*-butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]-N-ethylbenzenesulfonamide;*

15 *N-(4-{[[2-*tert*-butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl](ethyl)amino}sulfonyl)phenyl]-2,2-dimethylpropanamide;*

*2-[(4-{[[2-*tert*-butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl](ethyl)amino}sulfonyl)phenyl]amino]-2-oxoethyl acetate;*

*N-(4-{[[2-*tert*-butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl](ethyl)amino}sulfonyl)phenyl]-2-hydroxyacetamide;*

20 *N-[2-*tert*-butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]-N-ethyl-4-{{(isopropylamino)carbonyl}amino}benzenesulfonamide;*

*N-[4-({ethyl[2-(1-methoxy-1-methylethyl)-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]amino}sulfonyl)phenyl]acetamide;*

*4-[(aminocarbonyl)amino]-N-ethyl-N-[2-(1-methoxy-1-methylethyl)-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]benzenesulfonamide;*

25 *N-ethyl-N-[2-(1-methoxy-1-methylethyl)-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]-4-{{(methylamino)carbonyl}amino}benzenesulfonamide;*

*4-amino-N-ethyl-N-[2-(1-methoxy-1-methylethyl)-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]benzenesulfonamide;*

*N-[4-({ethyl[2-(1-methoxy-1-methylethyl)-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]amino}sulfonyl)phenyl]-2,2-dimethylpropanamide;*

30 *2-{{[4-({ethyl[2-(1-methoxy-1-methylethyl)-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]amino}sulfonyl)phenyl]amino}-2-oxoethyl acetate;*

N-[4-(*{ethyl}[2-(1-methoxy-1-methylethyl)-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]amino}sulfonyl)phenyl]-2-hydroxyacetamide;
N-ethyl-4-*{[(isopropylamino)carbonyl]amino}*-*N*-[2-(1-methoxy-1-methylethyl)-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]benzenesulfonamide;
5 *N*-(4-*{[[2-(1-methoxy-1-methylethyl)-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino}sulfonyl)phenyl)acetamide;
4-[(aminocarbonyl)amino]-*N*-[2-(1-methoxy-1-methylethyl)-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]-*N*-methylbenzenesulfonamide;
10 2-Hydroxy-*N*-(4-*{[[2-(1-methoxy-1-methylethyl)-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino}sulfonyl)phenyl)acetamide;
 N-(4-*{[[2-(1-ethoxy-1-methylethyl)-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino}sulfonyl)phenyl)acetamide;
 N-[4-(*{[1-(2-azetidin-1-ylethyl)-2-tert-butyl-1*H*-benzimidazol-5-yl]amino}sulfonyl)phenyl]acetamide;
15 3-[5-*{[4-(acetylamino)phenyl}sulfonyl]amino)-2-*tert*-butyl-1*H*-benzimidazol-1-yl]propyl acetate;
 N-{4-*{[(1S,4S)-bicyclo[2.2.1]hept-5-en-2-ylmethyl]-2-tert-butyl-1*H*-benzimidazol-5-yl}amino}sulfonyl)phenyl]acetamide;
 N-[4-*{[[2-tert-butyl-1-(tetrahydro-2*H*-pyran-3-ylmethyl)-1*H*-benzimidazol-5-yl]amino}sulfonyl)phenyl]acetamide;
20 *N*-{4-*{[(2-tert-butyl-1-[2-(tetrahydro-2*H*-pyran-4-yl)ethyl]-1*H*-benzimidazol-5-yl}amino}sulfonyl)phenyl]acetamide;
 N-{4-*{[(2-tert-butyl-1-(cyclobutylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino}sulfonyl)phenyl]acetamide;
25 4-[(aminocarbonyl)amino]-*N*-[2-*tert*-butyl-1-(cyclobutylmethyl)-1*H*-benzimidazol-5-yl]-*N*-methylbenzenesulfonamide;
 N-{4-*{[(2-tert-butyl-1-(cyclobutylmethyl)-1*H*-benzimidazol-5-yl)(methyl)amino}sulfonyl)phenyl]-2,2-dimethylpropanamide;
 N-(4-*{[(2-(1,1-difluoroethyl)-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino}sulfonyl)phenyl)-2-hydroxyacetamide;
30 *N*-(4-*{[(2-(1,1-difluoroethyl)-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino}sulfonyl)phenyl)acetamide;*************

N-(4-{[[2-(1,1-difluoroethyl)-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)-3-methylbutanamide;
5 *N*-(4-{[[2-(1,1-difluoroethyl)-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl](methyl)amino]sulfonyl}phenyl)-2,2-dimethylpropanamide;
N-[2-(1,1-difluoroethyl)-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-
10 yl]-4-{{(isopropylamino)carbonyl]amino}-*N*-methylbenzenesulfonamide;
4-{Bis[(isopropylamino)carbonyl]amino}-*N*-[2-(1,1-difluoroethyl)-1-(tetrahydro-
2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]-*N*-methylbenzenesulfonamide;
N-[4-({methyl}[1-(tetrahydro-2*H*-pyran-4-ylmethyl)-2-(trifluoromethyl)-1*H*-
15 benzimidazol-5-yl]amino]sulfonyl]phenyl]acetamide;
4-[(aminocarbonyl)amino]-*N*-methyl-*N*-[1-(tetrahydro-2*H*-pyran-4-ylmethyl)-2-
(trifluoromethyl)-1*H*-benzimidazol-5-yl]benzenesulfonamide;
N-methyl-4-nitro-*N*-[1-(tetrahydro-2*H*-pyran-4-ylmethyl)-2-(trifluoromethyl)-1*H*-
20 benzimidazol-5-yl]benzenesulfonamide;
4-amino-*N*-methyl-*N*-[1-(tetrahydro-2*H*-pyran-4-ylmethyl)-2-(trifluoromethyl)-
1*H*-benzimidazol-5-yl]benzenesulfonamide;
2,2-dimethyl-*N*-[4-({methyl}[1-(tetrahydro-2*H*-pyran-4-ylmethyl)-2-
25 (trifluoromethyl)-1*H*-benzimidazol-5-yl]amino]sulfonyl)phenyl]propanamide;
2-{{[4-({methyl}[1-(tetrahydro-2*H*-pyran-4-ylmethyl)-2-(trifluoromethyl)-1*H*-
benzimidazol-5-yl]amino]sulfonyl]phenyl]amino}-2-oxoethyl acetate;
4-{{(isopropylamino)carbonyl]amino}-*N*-methyl-*N*-[1-(tetrahydro-2*H*-pyran-4-
ylmethyl)-2-(trifluoromethyl)-1*H*-benzimidazol-5-yl]benzenesulfonamide;
2-Hydroxy-*N*-[4-({methyl}[1-(tetrahydro-2*H*-pyran-4-ylmethyl)-2-
(trifluoromethyl)-1*H*-benzimidazol-5-yl]amino]sulfonyl)phenyl]acetamide
and pharmaceutically acceptable salts thereof.

6. A compound according to any one of claims 1-5 for use as a medicament.

7. The use of a compound according to any one of claims 1-5 in the manufacture
30 of a medicament for the therapy of pain.

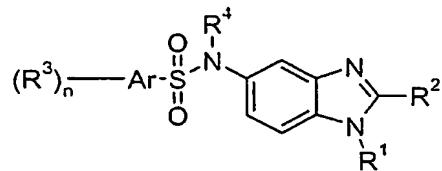
8. The use of a compound according to any one of claims 1-5 in the manufacture
of a medicament for the treatment of anxiety disorders.

9. The use of a compound according to any one of claims 1-5 in the manufacture of a medicament for the treatment of cancer, multiple sclerosis, Parkinson's disease, cancer, Huntington's chorea, Alzheimer's disease, gastrointestinal disorders and 5 cardiovascular disorders.

10. A pharmaceutical composition comprising a compound according to any one of claims 1-5 and a pharmaceutically acceptable carrier.

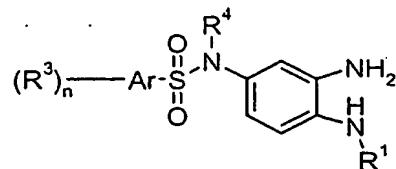
10 11. A method for the therapy of pain in a warm-blooded animal, comprising the step of administering to said animal in need of such therapy a therapeutically effective amount of a compound according to any one of claims 1-5.

12. A method for preparing a compound of Formula I,



15

comprising the step of reacting a compound of Formula II,



20 with a compound of $\text{R}^2\text{C}(=\text{O})\text{X}$, in the presence of a base and optionally a coupling reagent, followed by treatment with an acid; wherein

X is selected from Cl, Br, F and OH;

R^1 is selected from $\text{C}_{1-10}\text{alkyl}$, $\text{C}_{2-10}\text{alkenyl}$, $\text{C}_{2-10}\text{alkynyl}$, $\text{R}^5\text{C}(=\text{O})-\text{O}-$

25 $\text{C}_{1-6}\text{alkyl}$, $\text{R}^5\text{R}^6\text{N}-\text{C}_{1-6}\text{alkyl}$, $\text{R}^5\text{O}-\text{C}_{1-6}\text{alkyl}$, $\text{R}^5\text{C}(=\text{O})\text{N}(-\text{R}^6)-\text{C}_{1-6}\text{alkyl}$, $\text{R}^5\text{R}^6\text{NS}(=\text{O})_2-$ $\text{C}_{1-6}\text{alkyl}$, $\text{R}^5\text{CS}(=\text{O})_2\text{N}(-\text{R}^6)-\text{C}_{1-6}\text{alkyl}$, $\text{R}^5\text{R}^6\text{NC}(=\text{O})\text{N}(-\text{R}^7)-\text{C}_{1-6}\text{alkyl}$,

$R^5R^6NS(=O)_2N(R^7)-C_{1-6}alkyl$, $C_{6-10}aryl-C_{1-6}alkyl$, $C_{6-10}aryl-C(=O)-C_{1-6}alkyl$, $C_{3-10}cycloalkyl-C_{1-6}alkyl$, $C_{4-8}cycloalkenyl-C_{1-6}alkyl$, $C_{3-6}heterocyclyl-C_{1-6}alkyl$, $C_{3-6}heterocyclyl-C(=O)-C_{1-6}alkyl$, $C_{1-10}hydrocarbylamino$, R^5R^6N- , R^5O- , $R^5C(=O)N(-R^6)-$, $R^5R^6NS(=O)_2-$, $R^5CS(=O)_2N(-R^6)-$, $R^5R^6NC(=O)N(-R^7)-$, $R^5R^6NS(=O)_2N(R^7)-$,

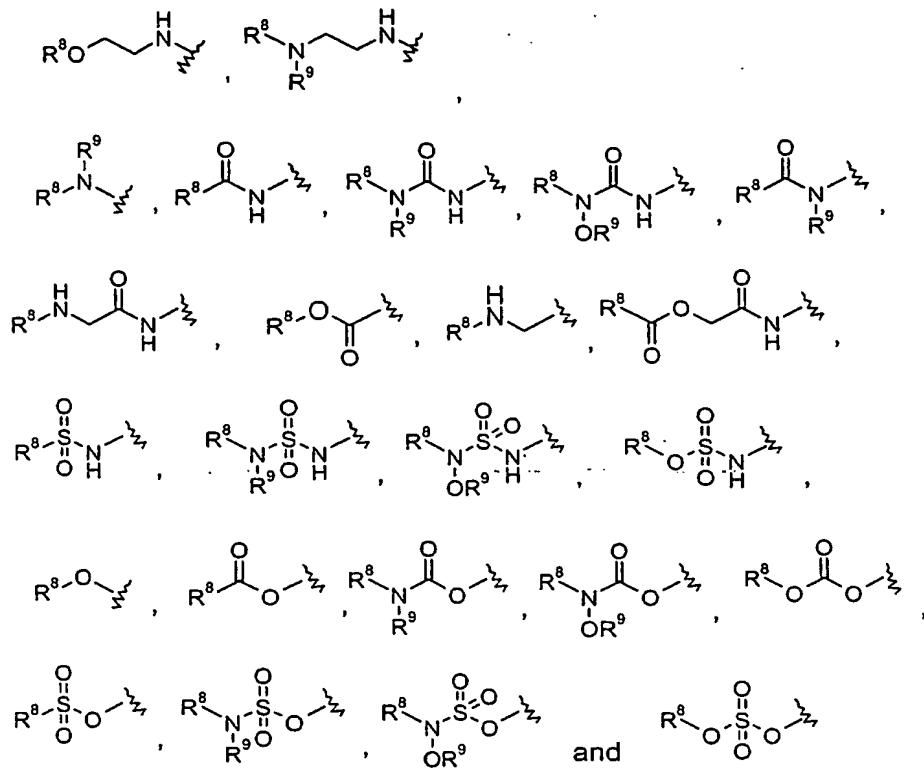
5 $C_{6-10}aryl$, $C_{6-10}aryl-C(=O)-$, $C_{3-10}cycloalkyl$, $C_{4-8}cycloalkenyl$, $C_{3-6}heterocyclyl$ and $C_{3-6}heterocyclyl-C(=O)-$; wherein said $C_{1-10}alkyl$, $C_{2-10}alkenyl$, $C_{2-10}alkynyl$, $C_{6-10}aryl-C_{1-6}alkyl$, $C_{6-10}aryl-C(=O)-C_{1-6}alkyl$, $C_{3-10}cycloalkyl-C_{1-6}alkyl$, $C_{4-8}cycloalkenyl-C_{1-6}alkyl$, $C_{3-6}heterocyclyl-C(=O)-C_{1-6}alkyl$, $C_{1-10}hydrocarbylamino$, $C_{6-10}aryl$, $C_{6-10}aryl-C(=O)-$, $C_{3-10}cycloalkyl$, $C_{4-8}cycloalkenyl$, $C_{3-6}heterocyclyl$ or $C_{3-6}heterocyclyl-C(=O)-$ used in defining R^1 is optionally substituted by one or more groups selected from halogen, cyano, nitro, methoxy, ethoxy, methyl, ethyl, hydroxy, benzyl, and $-NR^5R^6$;

10 R^2 is selected from $C_{1-10}alkyl$, $C_{2-10}alkenyl$, $C_{2-10}alkynyl$, $C_{3-10}cycloalkyl$, $C_{3-10}cycloalkyl-C_{1-6}alkyl$, $C_{4-8}cycloalkenyl-C_{1-6}alkyl$, $C_{3-6}heterocycloalkyl-C_{1-6}alkyl$, $C_{4-8}cycloalkenyl$, R^5R^6N- , $C_{3-5}heteroaryl$, $C_{6-10}aryl$ and $C_{3-6}heterocycloalkyl$, wherein said $C_{1-10}alkyl$, $C_{2-10}alkenyl$, $C_{2-10}alkynyl$, $C_{3-8}cycloalkyl$, $C_{3-8}cycloalkyl-C_{1-6}alkyl$, $C_{4-8}cycloalkenyl-C_{1-6}alkyl$, $C_{3-6}heterocycloalkyl-C_{1-6}alkyl$, $C_{4-8}cycloalkenyl$, $C_{3-6}heteroaryl$, $C_{6-10}aryl$ or $C_{3-6}heterocycloalkyl$ used in defining R^2 is optionally substituted by one or more groups selected from halogen, cyano, nitro, methoxy, ethoxy, methyl, ethyl, hydroxy, and $-NR^5R^6$;

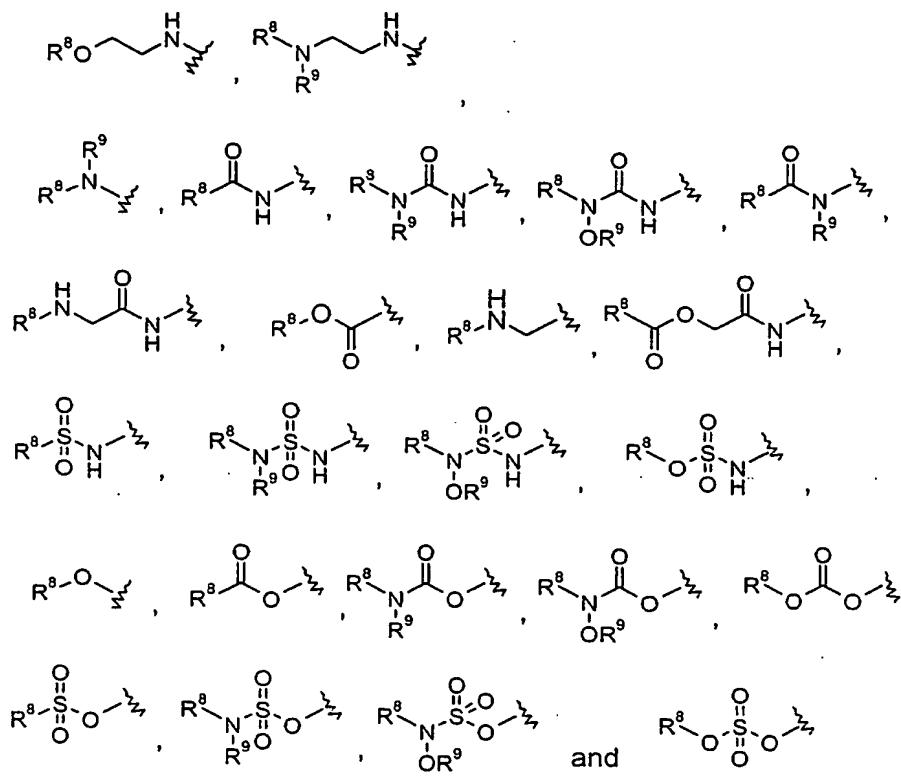
15 wherein R^5 , R^6 and R^7 are independently selected from $-H$, $C_{1-6}alkyl$, $C_{2-6}alkenyl$, $C_{2-6}alkynyl$, and a divalent C_{1-6} group that together with another divalent R^5 , R^6 or R^7 forms a portion of a ring;

20 Ar is selected from $C_{6-10}aryl$ and $C_{3-8}heteroaryl$;

25 n is selected from 0, 1, 2 and 3; each of R^3 is independently selected from $-H$, nitro, halogen, $C_{1-10}alkyl$, $C_{2-10}alkenyl$, $C_{2-10}alkynyl$, $C_{3-10}cycloalkyl$, $C_{3-10}cycloalkyl-C_{1-6}alkyl$, $C_{4-8}cycloalkenyl-C_{1-6}alkyl$, $C_{3-6}heterocycloalkyl-C_{1-6}alkyl$, $C_{3-6}heterocycloalkyl$



optionally substituted with one or more groups selected from C₁₋₆alkyl, hydroxy, halogen, amino and C₁₋₆alkoxy,



each of R⁸ and R⁹ is independently selected from -H, C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₆alkyl, C₃₋₆heterocyclyl, C₆₋₁₀aryl, C₃₋₆heterocyclyl-C₁₋₆alkyl, C₆₋₁₀aryl-C₁₋₆alkyl, and a divalent C₁₋₆group that together
 5 with another divalent group selected from R⁸ and R⁹ forms a portion of a ring, wherein said C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₆alkyl, C₃₋₆heterocyclyl, C₆₋₁₀aryl, C₃₋₆heterocyclyl-C₁₋₆alkyl, C₆₋₁₀aryl-C₁₋₆alkyl, or divalent C₁₋₆group is optionally substituted by one or more groups selected from halogen, cyano, nitro, methoxy, ethoxy, methyl, ethyl, hydroxy, and -NR⁵R⁶; and
 10 R⁴ is selected from -H, C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₆alkyl, and C₄₋₈cycloalkenyl-C₁₋₆alkyl.